

20090309-93444007

The new substrate is synthetically easily accessible

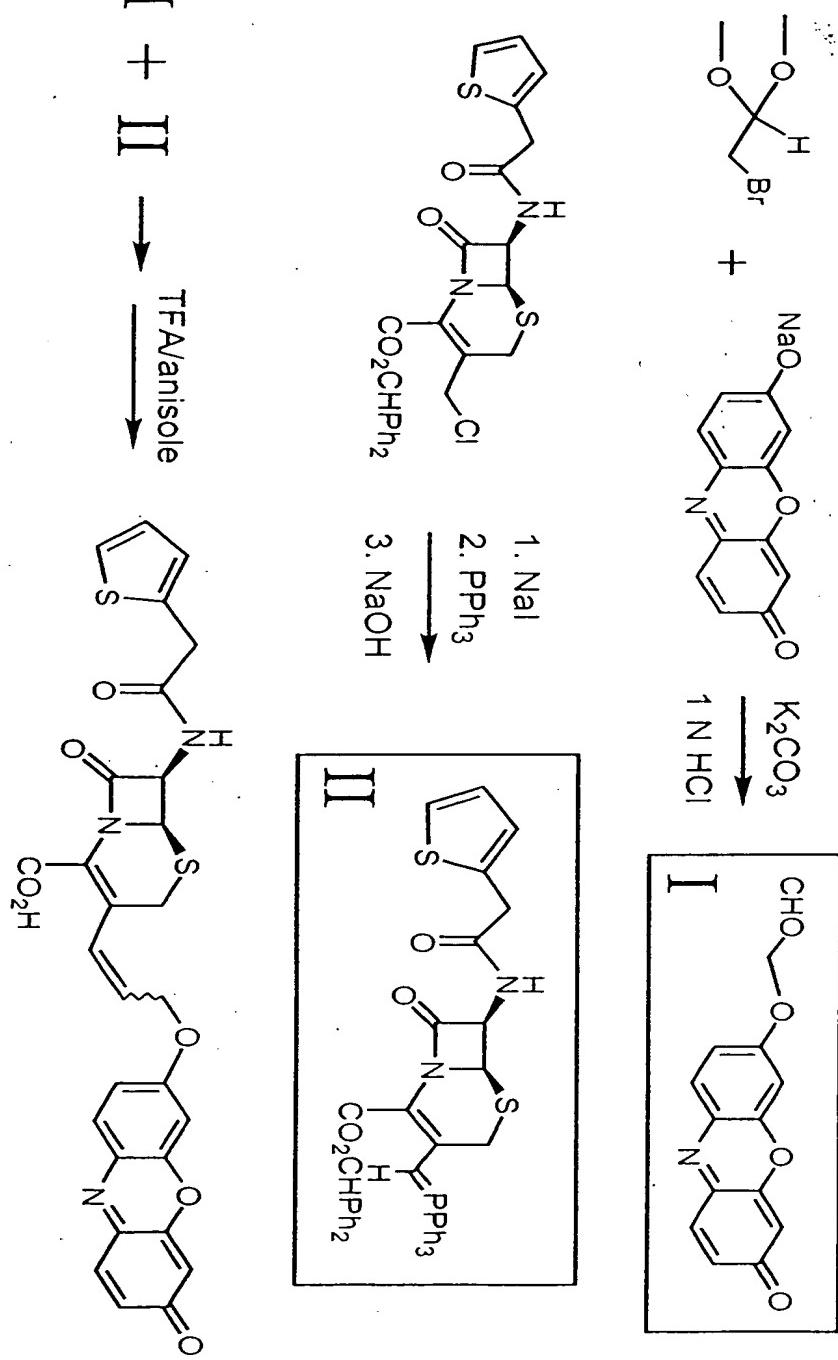


FIG. 1

Enzymatic fragmentation can take place to the new substrate

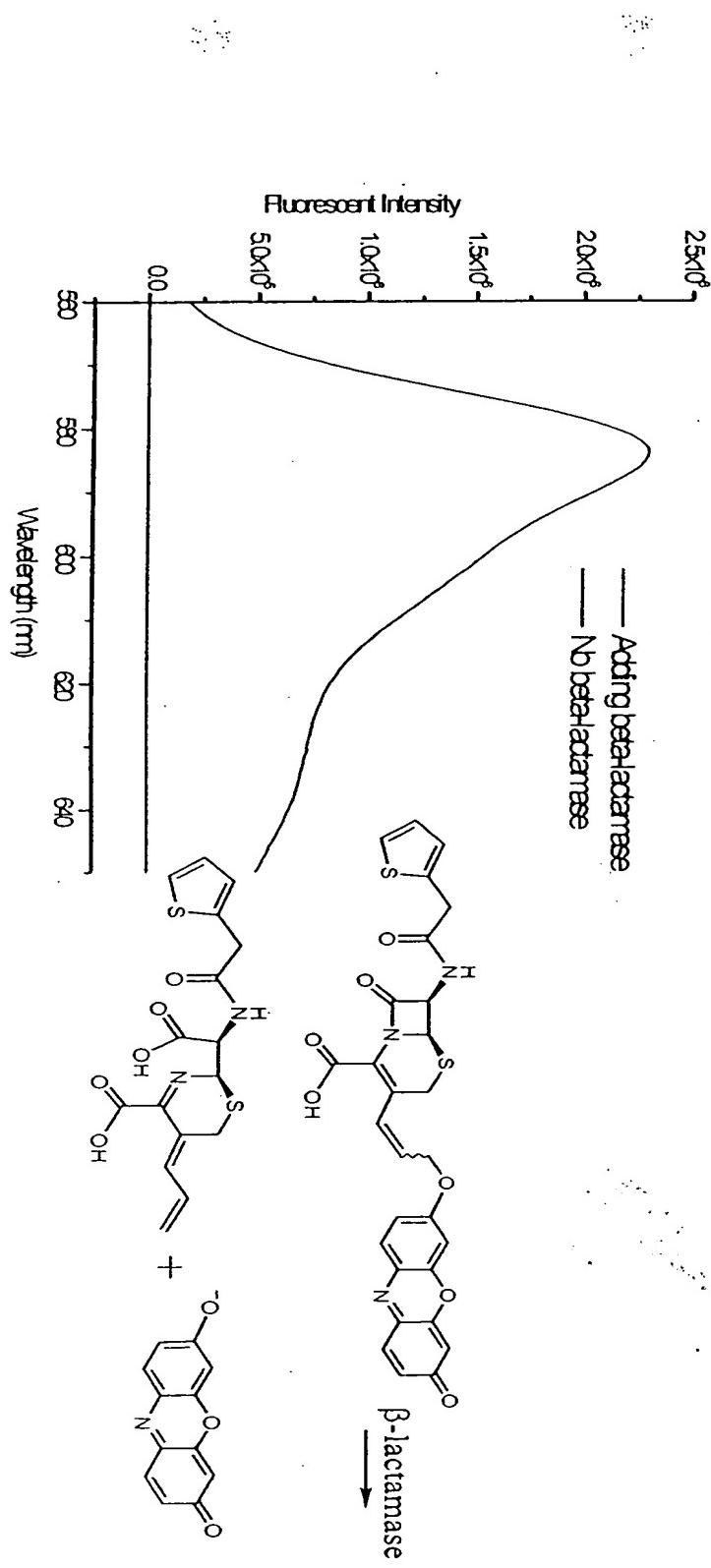


FIG 2

20200907 - 33444000 T

Synthesis of RECTO

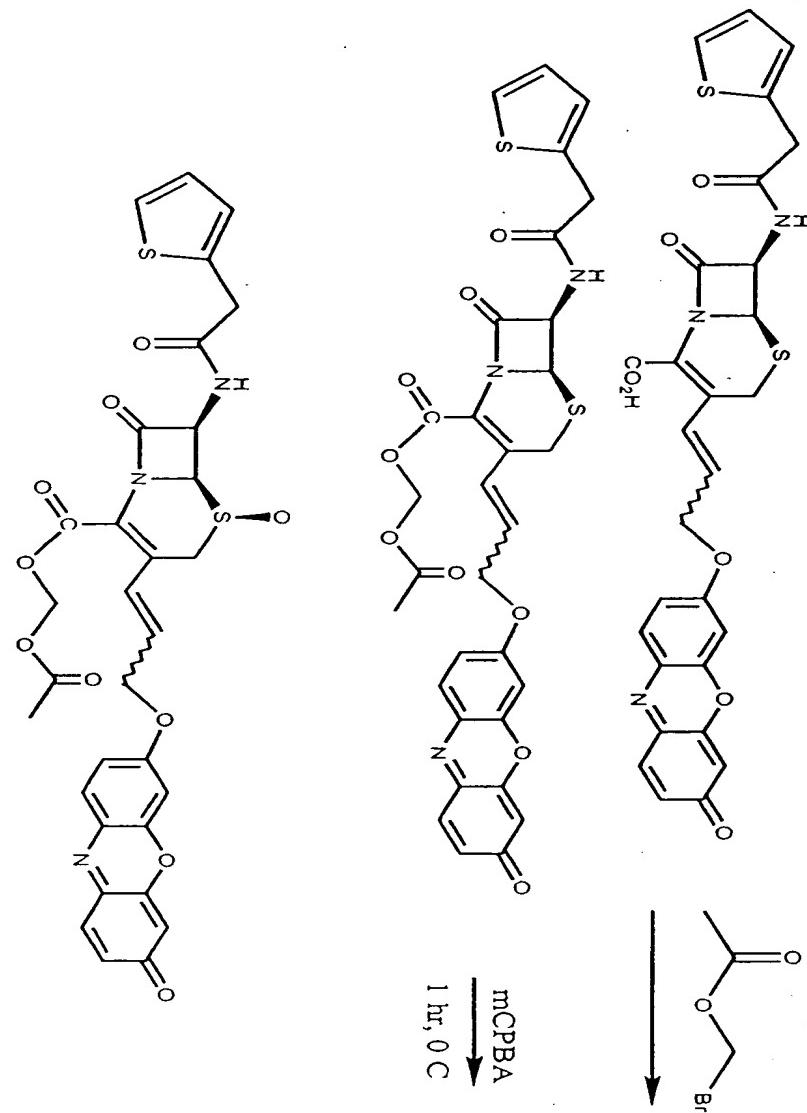


FIG. 3

Oxidation state of the sulfide affects stability of the substrate

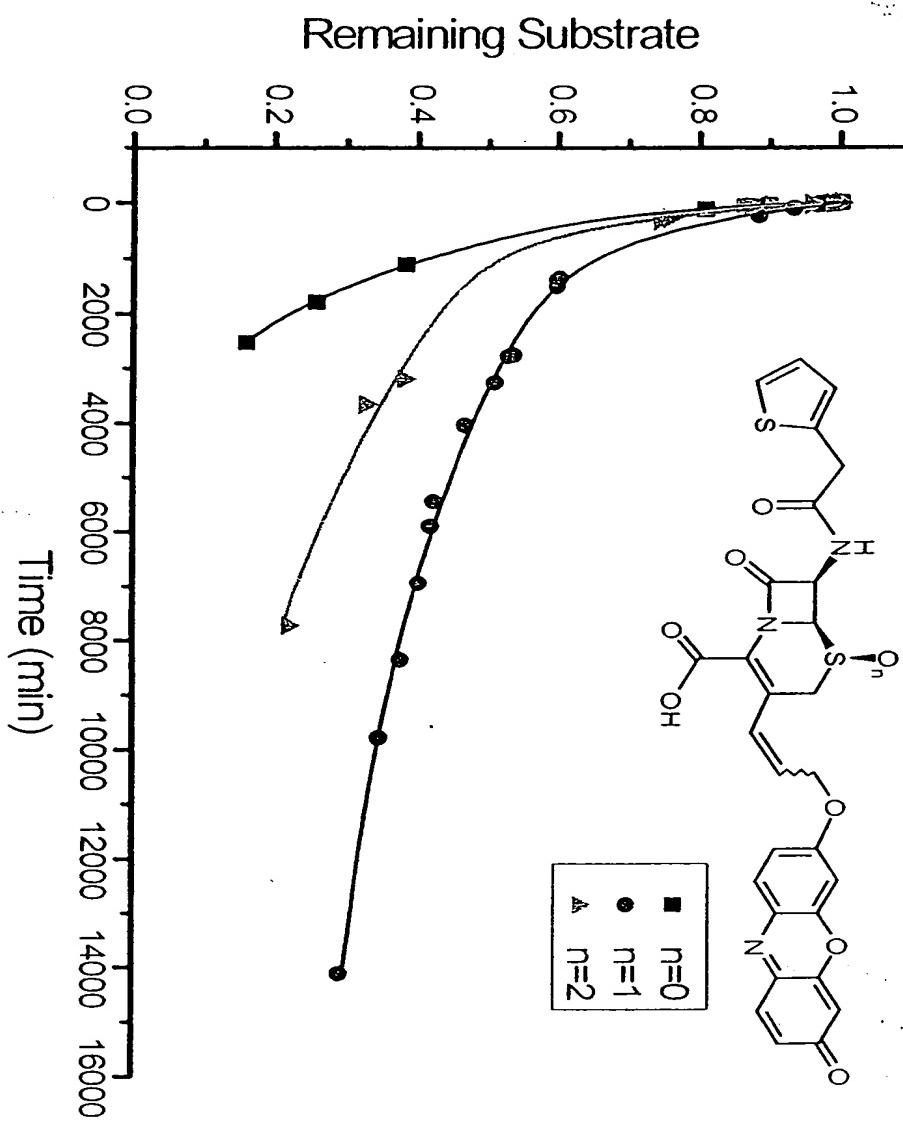


FIG 4

Sulfoxide increases substrate stability

20090 - 5844000 T

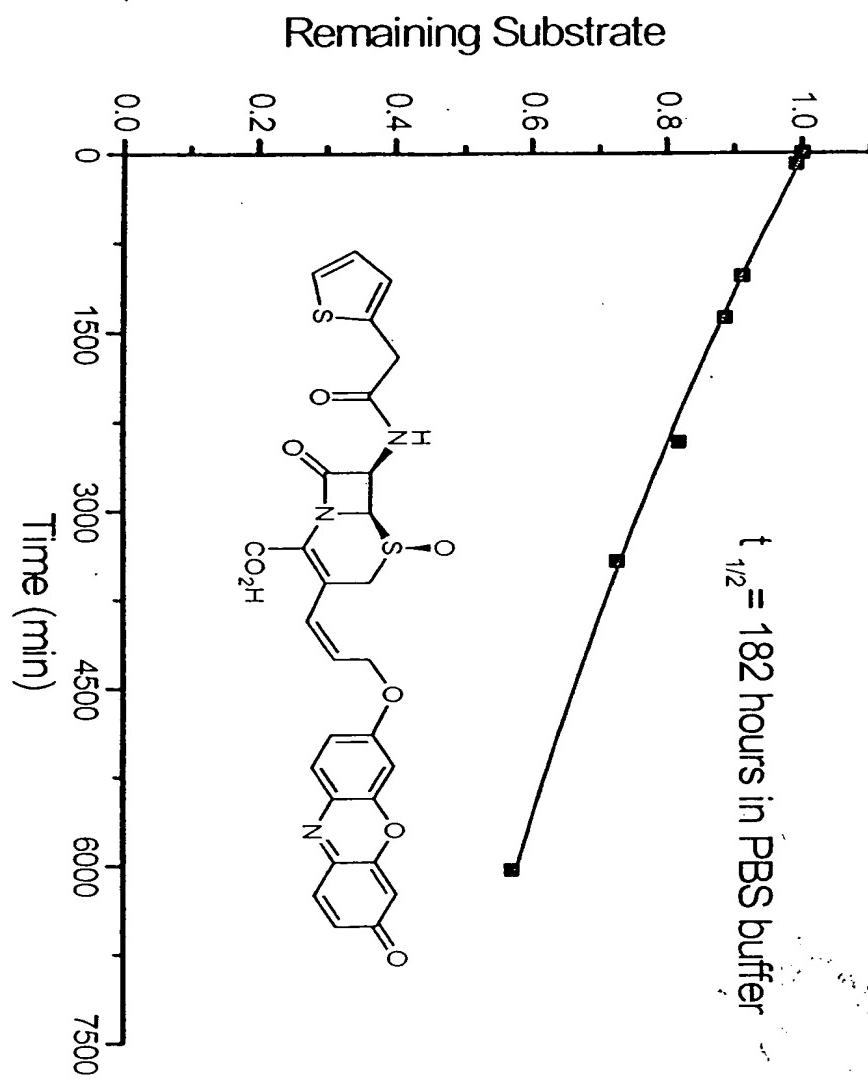
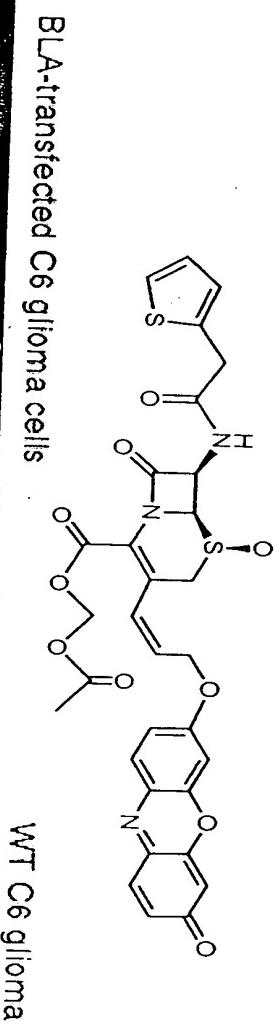


FIG. 5

203050 " 33444400 T

Increased resorufin deposition in β -lactamase-transfected vs. wild type cells



β LA-transfected C6 glioma cells

WT C6 glioma

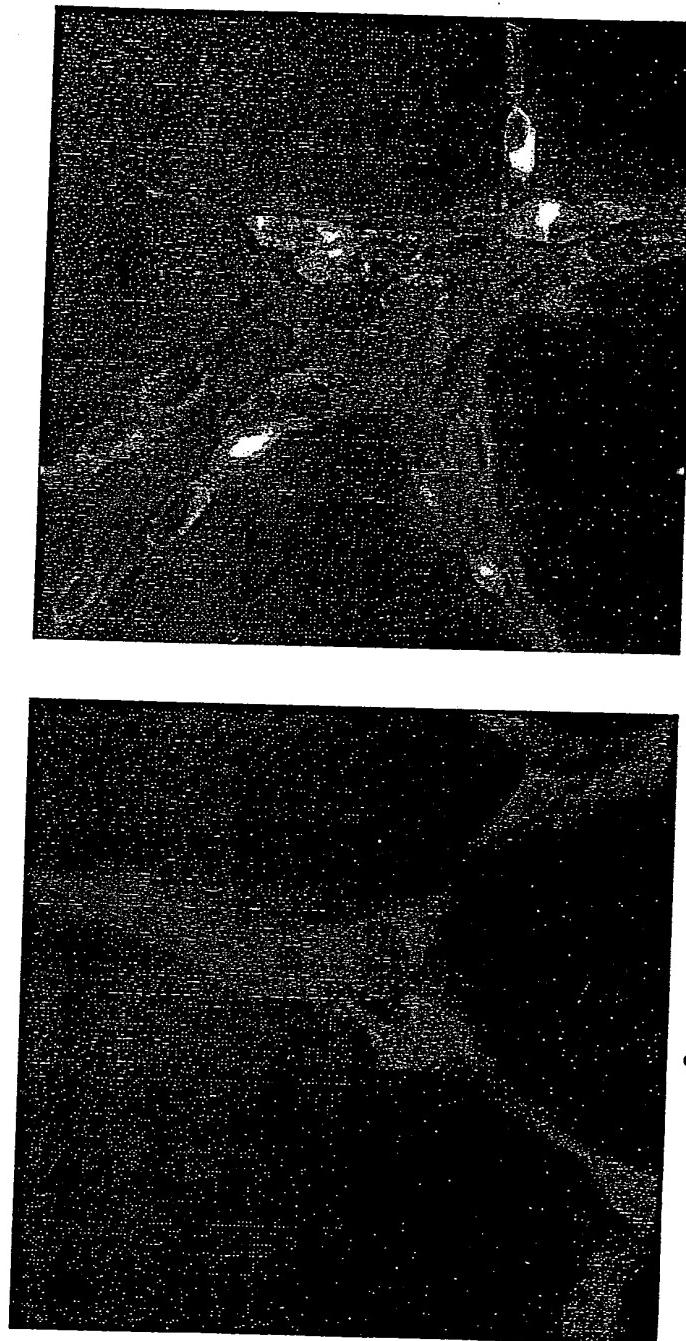
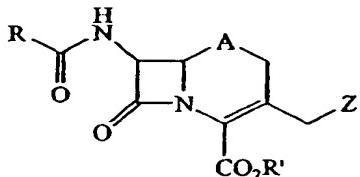


FIG. 6

cephalosporin-phenol ethers that we wish to claim:



Preferred R = benzyl, 2-thienylmethyl, or cyanomethyl; A = S or SO; R' = H or physiologically acceptable salts or ester groups.

where Z can be:

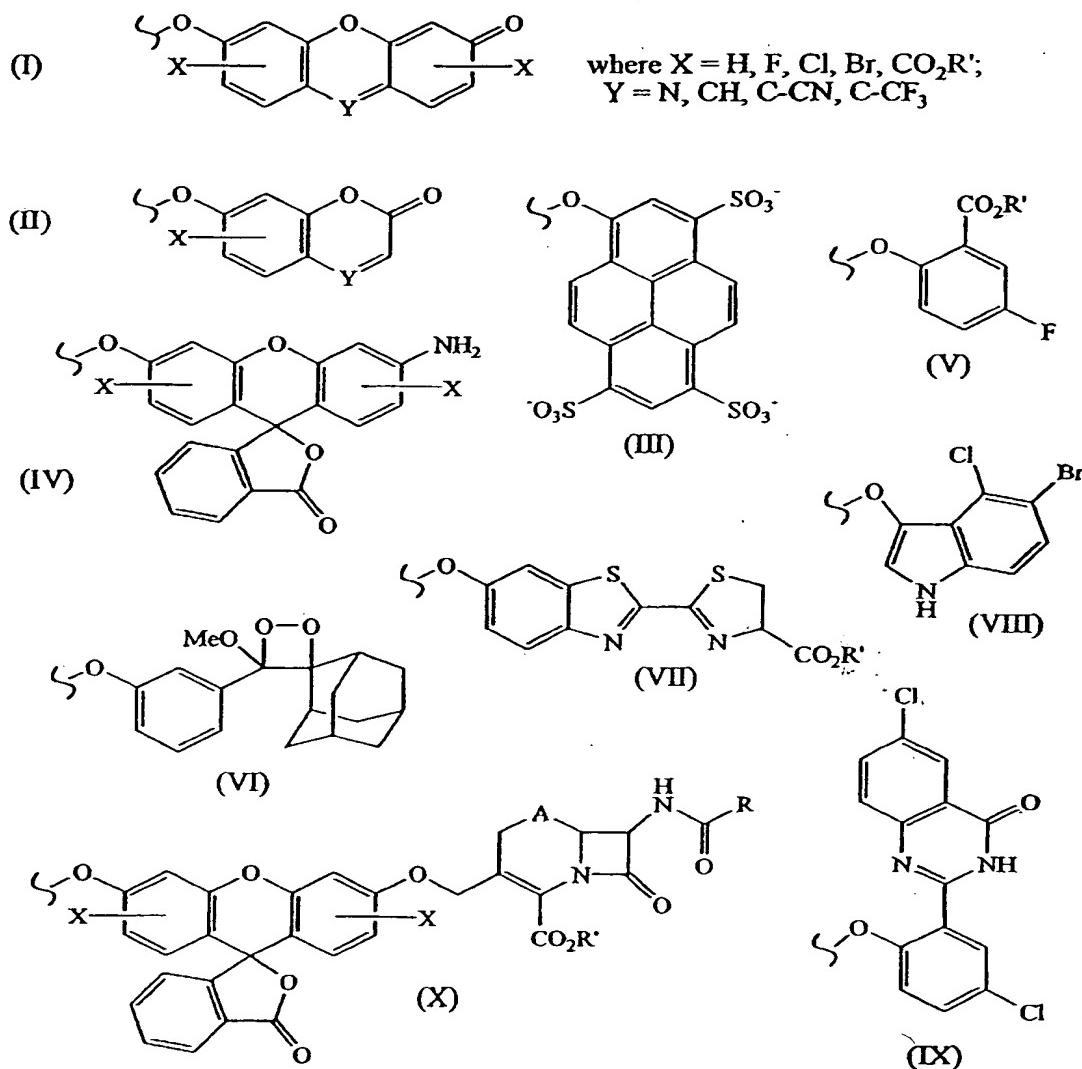


FIG 7

Resorufin-cephalosporin cleaved by β -lactamase

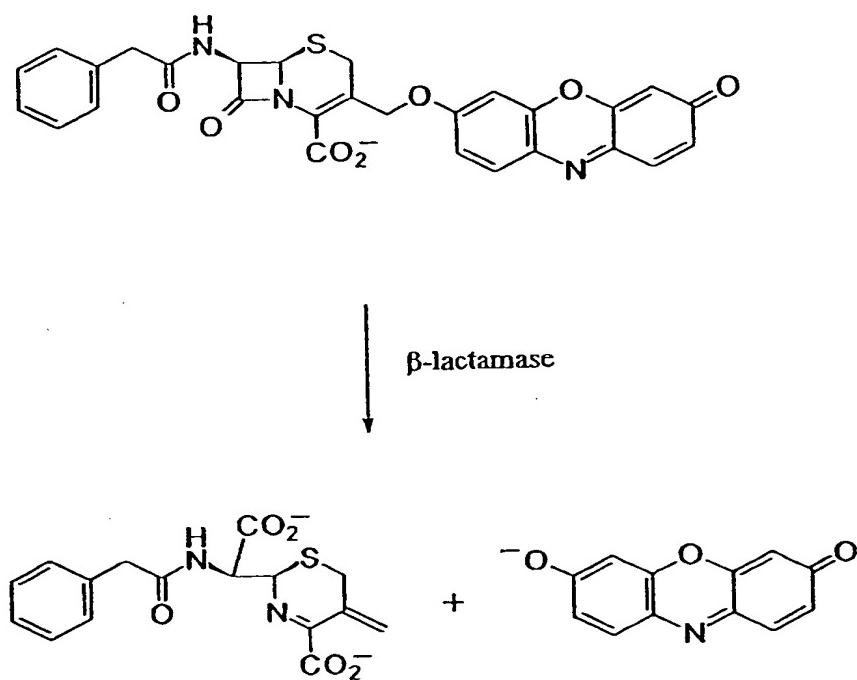


FIG 8

203030 - 98444000 T

Absorption spectra of resorufin-cephalosporin

before and after β -lactamase treatment

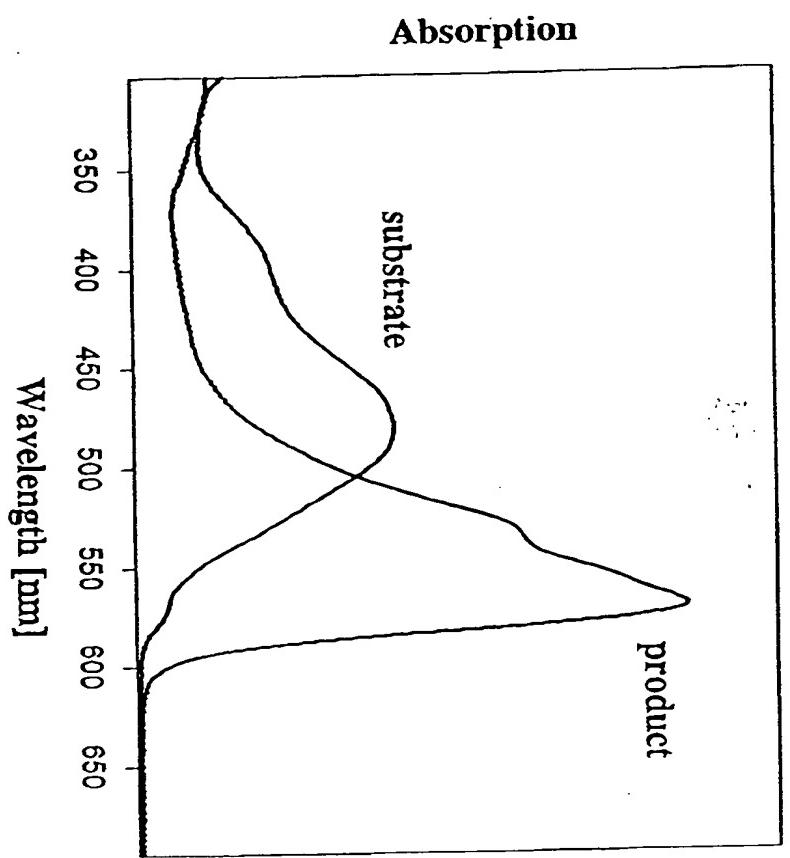


FIG. 9

203090 " 36414100 T

Fluorescence emission of resorufin-cephalosporin
before and after β -lactamase treatment

(excitation at 570 nm)

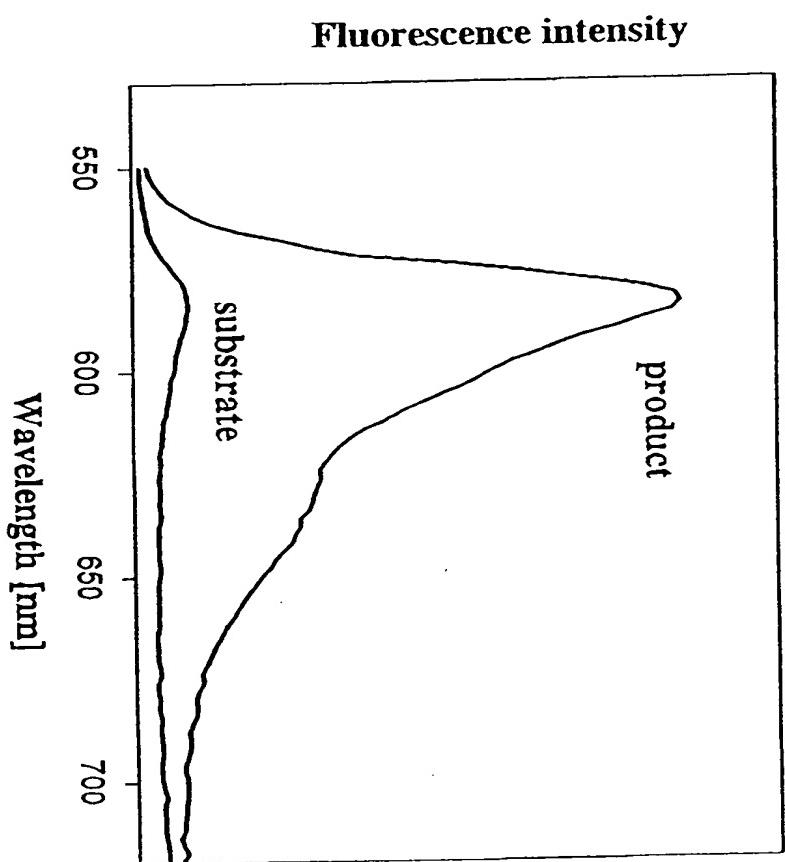


FIG. 10